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| APPLICATION NO. | FILING DATE | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. | CONFIRMATION NO. |
|----------------------|---------------------------------------|----------------------|----------------------------|------------------|
| 10/573,378 | 03/24/2006 | Kenya Shitara | 00005.01290 | 4546 |
| 5514 FITZPATRICI | 7590 08/16/2007 K CELLA HARPER & S | | EXAMINER | |
| 30 ROCKEFELLER PLAZA | | | DANG, IAN D | |
| NEW YORK, | NY 10112 | | ART UNIT PAPER NUMBER 1647 | |
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| | | | MAIL DATE | DELIVERY MODE |
| | | | 08/16/2007 | PAPER |

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

| | | Application No. | Applicant(s) | | | |
|--|--|--|---|--|--|--|
| Office Action Summary | | 10/573,378 | SHITARA ET AL. | | | |
| | | Examiner | Art Unit | | | |
| • | • | lan Dang | 1647 | | | |
| The MAILIN Period for Reply | The MAILING DATE of this communication appears on the cover sheet with the correspondence address | | | | | |
| A SHORTENED S WHICHEVER IS L - Extensions of time may after SIX (6) MONTHS I - If NO period for reply is - Failure to reply within th Any reply received by th | TATUTORY PERIOD FOR REPLY ONGER, FROM THE MAILING DA be available under the provisions of 37 CFR 1.13 from the mailing date of this communication. specified above, the maximum statutory period we set or extended period for reply will, by statute, the Office later than three months after the mailing stment. See 37 CFR 1.704(b). | ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tire will apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE | N. nely filed the mailing date of this communication. ED (35 U.S.C. § 133). | | | |
| Status | | | | | | |
| 2a) ☐ This action is 3) ☐ Since this ap | to communication(s) filed on s FINAL 2b)⊠ This oplication is in condition for allowar cordance with the practice under E | action is non-final. nce except for formal matters, pro | | | | |
| Disposition of Claims | 3 | | | | | |
| 4a) Of the ab 5) ☐ Claim(s) 6) ☐ Claim(s) 7) ☐ Claim(s) | | vn from consideration. | | | | |
| Application Papers | | | | | | |
| 9) The specification is objected to by the Examiner. 10) The drawing(s) filed on is/are: a) accepted or b) objected to by the Examiner. Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a). Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d). 11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152. | | | | | | |
| Priority under 35 U.S | .C. § 119 | | • | | | |
| 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. | | | | | | |
| | | | | | | |
| · <u> </u> | n's Patent Drawing Review (PTO-948) e Statement(s) (PTO/SB/08) | 4) Interview Summary Paper No(s)/Mail D 5) Notice of Informal I 6) Other: | ate | | | |

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DETAILED ACTION

Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 1-21, 29 and 30 drawn to a recombinant antibody.

Group II, claim(s) 22-25 drawn to DNA encoding the recombinant antibody and a method for producing the antibody.

Group III, claim(s) 27-28, drawn to a method of treat IGF- associated diseases, which comprises administering a therapeutically effective amount of the recombinant or the antibody fragment.

The inventions listed as Groups I-II do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

According to PCT Rule 13.2, unity of invention exists only when the shared same or corresponding technical feature is a contribution over the prior art. The inventions listed as Groups I-IV do not relate to a single general inventive concept because they lack the same or corresponding technical feature.

Claim 1 is directed to a recombinant antibody or an antibody fragment wherein the recombinant antibody or the antibody fragment binds to human insulin-like growth factor-I (IGF-I) and human insulin-like growth factor-II (IGF-II) to inhibit the biological activities of human IGF-I and human IGF-II. Cohen al. (WO 02/053596 A2, filed 12/20/2001, and published 07/11/2002)

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teach a recombinant antibody or an antibody fragment wherein the recombinant antibody or the antibody fragment binds to human insulin-like growth factor-I (IGF-I) and human insulin-like growth factor-II (IGF-II) to inhibit the biological activities of human IGF-I and human IGF-II (page 89, claims 1-3). The prior art meets the limitations disclosed in claim 1. Thus Group I lacks novelty or inventive step and does not make a contribution over the prior art. Since the first claimed invention has no special technical feature, it cannot share a special technical feature with the other claimed invention.

Under PCR Rule 13.1, the application shall relate to one invention only or to a group of inventions so linked as to form a single general inventive concept.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

(1) This application contains claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

The species for amino acid sequences encoding the VH of an antibody are as follows:

- a) CDR1, CDR2, CDR3 Antibody VH represented by SEQ ID Nos 5, 6, and 7
- b) SEQ ID NO:26

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Applicant is required, in reply to this action, to elect a single species to which the claims shall be restricted if no generic claim is finally held to be allowable. The reply must also identify the claims readable on the elected species, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered non-responsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

The claims are deemed to correspond to the species listed above in the following manner: claims 6 and 14.

The following claim(s) are generic: claim 5.

The species listed above do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, the species lack the same or corresponding special technical features for the following reasons: the sequences listed in claims 6 and 14 do not share a common structural feature because they have different sequences.

(2) This application contains claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

The species for amino acid sequences encoding the VL of an antibody are as follows:

- c) CDR1, CDR2, CDR3 Antibody VL represented by SEQ ID NOs: 8, 9, and 10
- d) SEQ ID NO:27
- e) SEQ ID NO:28

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f) SEQ ID NO:29

Applicant is required, in reply to this action, to elect a single species to which the claims shall be restricted if no generic claim is finally held to be allowable. The reply must also identify the claims readable on the elected species, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered non-responsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

The claims are deemed to correspond to the species listed above in the following manner: claims 7 and 15-19.

The following claim(s) are generic: claim 5.

The species listed above do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, the species lack the same or corresponding special technical features for the following reasons: the sequences listed in claims 7 and 15-19 do not share a common structural feature because they have different sequences.

Information

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Ian Dang whose telephone number is (571) 272-5014. The examiner can normally be reached on Monday-Friday from 9am to 5pm.

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If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Nickol can be reached on (571) 272-0835. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Ian Dang Patent Examiner Art Unit 1647 July 31st, 2007

> BRIDGET E. BUNNER PRIMARY EXAMINER

Didget P. Durner